INSTANTFOAM ALCOHOL HAND SANITIZER - ethyl alcohol liquid
Deb USA, Inc.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Drug Facts

Active Ingredient

Ethyl Alcohol 72%

Purpose

Antibacterial

Uses

For hand sanitizing to reduce bacteria on skin

Warnings

For external use only

Avoid contact with eyes

Flammable

Keep away from flame or fire

Keep out of reach of children.

Consult physician or poison control if ingested.

Directions

Apply one shot to dry hands

Rub into skin

No rinsing required

Inactive Ingredients

Water, n-Propanol, Cocoglucoside, Glyceryl Oleate, PEG-7 Glyceryl Cocoate, PEG-200 Hydrogenated Glyceryl Palmate, Dihydroxypropyl PEG-5 Linoleammonium Chloride, Behentrimonium Chloride, Bis-PEG-12 Dimethicone

deb

Instant Foam

alcohol hand sanitizer

refreshing

no water required

use anywhere, any time

use everyday

Kills>99.9999%

of common germs in 15 seconds

Made in Canada

Worldwide Patent Pending

deb foam technology

NSF

Nonfood Compounds Program Listed E-3 140062

01827-01-116

1 Liter - 33.8 Fluid Ounces

INSTANTFOAM ALCOHOL HAND SANITIZER

ethyl alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11084-017
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
Alcohol (UNII: 3K9958V90M) (Alcohol - UNII:3K9958V90M)	Alcohol	72 mL in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
Water (UNII: 059QF0KO0R)	
DIHYDROXYPROPYL PEG-5 LINOLEAMMONIUM CHLORIDE (UNII: 0Y0NQR2GH1)	
Propyl Alcohol (UNII: 96F264O9SV)	
Behentrimonium Chloride (UNII: X7GNG3S47T)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:11084-017-01	47 mL in 1 BOTTLE, PUMP		
2	NDC:11084-017-40	400 mL in 1 BOTTLE, SPRAY		
3	NDC:11084-017-27	1000 mL in 1 BOTTLE, SPRAY		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333	06/02/2008	

Labeler - Deb USA, Inc. (607378015)

Establishment

Name	Address	ID/FEI	Business Operations
Deb Worldwide Healthcare Inc.		205662831	manufacture

Revised: 8/2010 Deb USA, Inc.